

CLAIMS:

1) A method of imparting, controlling or improving the charge of an electrophotographic toner or developer, of a powder coating, of an electret material or in an electrostatic separation process, comprising the step of adding a salt-like structured silicate in which the cation is NH_4^+ , H_3O^+ , an alkali metal, alkaline earth metal, earth metal or transition metal ion or a low molecular weight organic cation or a combination thereof and the anion is an island, cyclic, group, chain, ribbon, laminar or matrix silicate or a combination thereof to a binder of an electrophotographic toner or developer or of a powder coating, to an electret fiber material or to an electrostatic separation process.

2) The method as claimed in claim 1, wherein the silicate is an anion selected from the group consisting of montmorillonite, bentonite, hectorite, kaolinite, serpentine, talc, pyrophyllite, mica, phlogopite, biotite, muscovite, paragonite, vermiculite, beidellite, xantophyllite, margarite, feldspar, zeolite, wollastonite, actinolite, amosite, crocidolite, sillimanite, nontronite, smectite, sepiolite, saponite, faujasite, permutite and sasil.

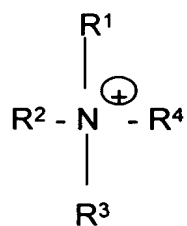
3) The method as claimed in claim 1, wherein the cation is H_3O^+ , Li^+ , Na^+ , K^+ , Rb^+ , Cs^+ , Be^{2+} , Mg^{2+} , Ca^{2+} , Sr^{2+} , Ba^{2+} , Al^{3+} , TiO^{2+} , ZrO^{2+} , Zn^{2+} , Fe^{2+} , Fe^{3+} , Sn^{2+} , Sn^{4+} , Pb^{2+} , Pb^{4+} , Cr^{3+} , Mn^{4+} , Mn^{2+} , Co^{2+} , Co^{3+} , Cu^{2+} , Sc^{3+} , Ti^{4+} , Zr^{4+} , V^{5+} , Y^{3+} , Ni^{2+} , Mo^{6+} or W^{6+} .

4) The method as claimed in claim 1, wherein the low molecular weight organic cation is a substituted ammonium, phosphonium, thionium or triphenylcarbonium ion or a cationic metal complex.

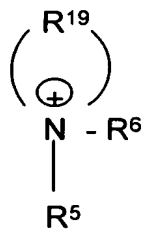
5) The method as claimed in claim 1, wherein the ammonium ion has one of the formulae (a) - (j)

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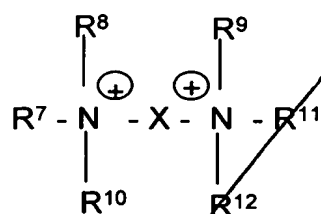
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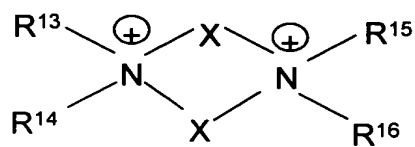
(a)



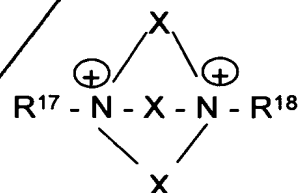
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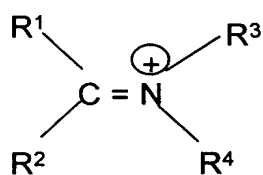
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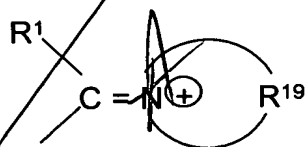
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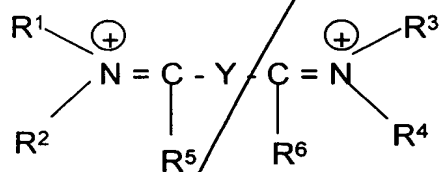
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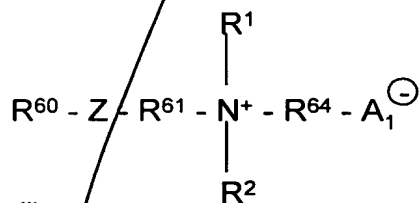
(f)



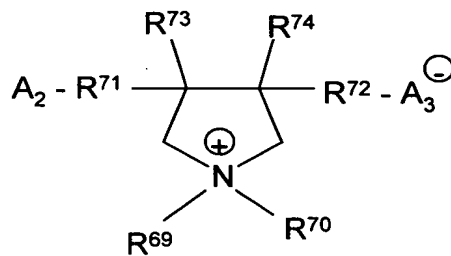
(g)



(h)



(i)



(j)

in which

R¹ to R¹⁸ are identical or different and represent hydrogen, CN, (CH₂)₁₋₁₈CN, halogen, branched or unbranched C₁-C₃₂-alkyl, mono- or polyunsaturated C₂-C₃₂-alkenyl, C₁-C₂₂-alkoxy, C₁-C₂₂-hydroxyalkyl, C₁-C₂₂-halogenoalkyl, C₂-C₂₂-halogenoalkenyl, C₁-C₂₂-aminoalkyl, (C₁-C₁₂)-trialkyl-ammonium-(C₁-C₂₂)-alkyl; (C₁-C₂₂)-alkylene-(C=O)O-(C₁-C₃₂)alkyl, (C₁-C₂₂)-alkylene-(C=O)O-aryl, (C₁-C₂₂)-alkylene-(C=O)NH-(C₁-C₃₂)alkyl, (C₁-C₂₂)-alkylene-(C=O)NH-aryl, (C₁-C₂₂)-alkylene-O(CO)-(C₁-C₃₂)alkyl, (C₁-C₂₂)-alkylene-O(CO)aryl, (C₁-C₂₂)-alkylene-NH(C=O)-(C₁-C₃₂)alkyl, (C₁-C₂₂)-alkylene-NHCO-aryl,

wherein



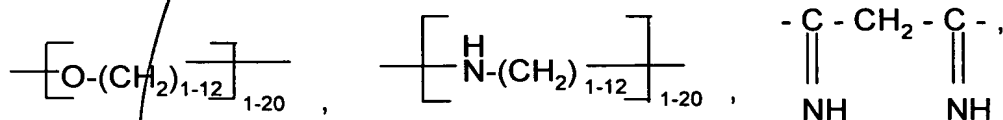
are optionally inserted into the acid ester or acid amide bonds;

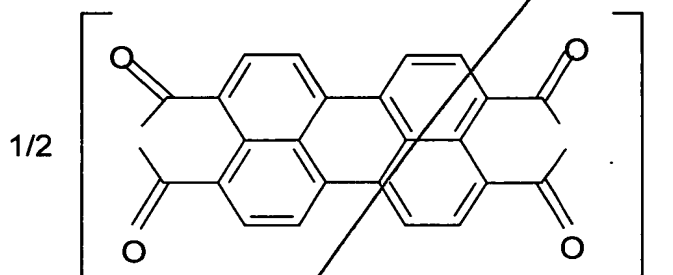
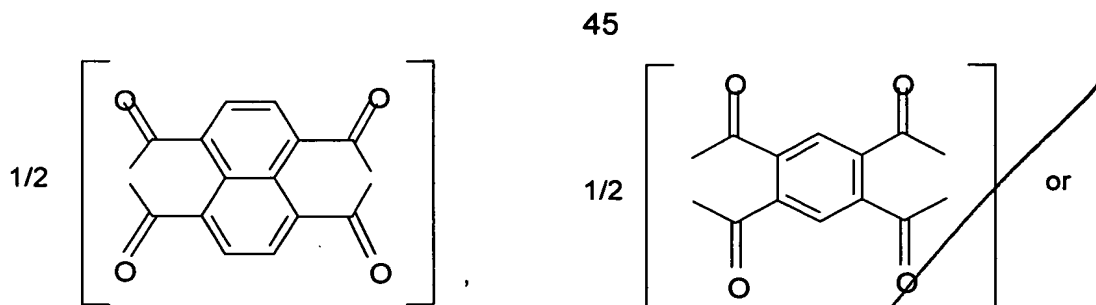
[(C₁-C₁₂)-alkylene-O]₁₋₁₀₀-H; aryl, (C₁-C₁₈)-alkylenearyl; -(O-SiR'₂)₁₋₃₂-O-SiR'₃, in which R' has the meaning C₁-C₁₂-alkyl, phenyl, benzyl or C₁-C₁₂-alkoxy;

heterocyclyl, C₁-C₁₈-alkylene-heterocyclyl, wherein the aryl and heterocyclyl radicals are optionally mono- or polysubstituted on carbon atoms or heteroatoms by C₁-C₁₂-alkyl, C₁-C₄-alkenyl, C₁-C₄-alkoxy, hydroxy-(C₁-C₄)alkyl, amino-(C₁-C₄)alkyl, C₁-C₄-alkylimino, carboxyl, hydroxyl, amino, nitro, cyano, halogen, C₁-C₁₂-acyl, C₁-C₄-halogenoalkyl, C₁-C₄-alkylcarbonyl, C₁-C₄-alkylcarbonyloxy, C₁-C₄-alkoxycarbonyl, C₁-C₄-alkylaminocarbonyl, C₁-C₄-alkylcarbonylimino, C₆-C₁₀-arylcarbonyl, aminocarbonyl, aminosulfonyl, C₁-C₄-alkylaminosulfonyl, phenyl, naphthyl, or heteroaryl,;

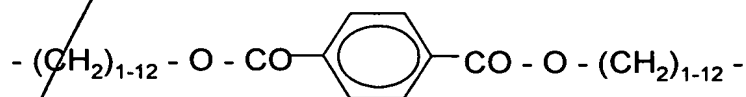
R¹⁹ represents C₄-C₁₁-alkylene, -(C₂H₄-O)₁₋₁₇-(CH₂)₁₋₂-, -(C₂H₄-NR-)₁₋₁₇-(CH₂)₁₋₂-, in which R is hydrogen or C₁-C₁₂-alkyl;

X has the meaning of Y and -CO-CH₂-CO-,





Y has the meaning $\text{-}\overset{\text{O}}{\underset{\text{||}}{\text{C}}}\text{-}$, $\text{-}\overset{\text{S}}{\underset{\text{||}}{\text{C}}}\text{-}$, $\text{-}\overset{\text{NH}}{\underset{\text{||}}{\text{C}}}\text{-}$, $\text{-(CH}_2\text{)}_{1-18}\text{-}$,



or o-, p-, m-(C₆-C₁₄)-arylene or (C₄-C₁₄)-heteroarylene with 1, 2, 3 or 4 heteroatoms from the group consisting of N, O, S or a combination thereof;

R⁶⁰ represents C₁-C₃₂-acyl, C₁-C₂₂-alkyl, C₂-C₂₂-alkenyl, C₁-C₁₈-alkylene-C₆-C₁₀-aryl, C₁-C₂₂-alkylene-heterocyclyl, C₆-C₁₀-aryl or (C₄-C₁₄)-heteroaryl with 1, 2, 3 or 4 heteroatoms from the group consisting of N, O, S, or a combination thereof;

R^{73} and R^{74} denote hydrogen or (C₁-C₁₈)-alkyl.

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- 7) The method as claimed in claim 4, wherein the ammonium ion is an aliphatic or aromatic 5- to 12-membered heterocyclic radical with 1 to 4 atoms selected from the group consisting of N, O and S, or a combination thereof, belonging to the rings, it being possible for 2 to 8 rings to be fused.

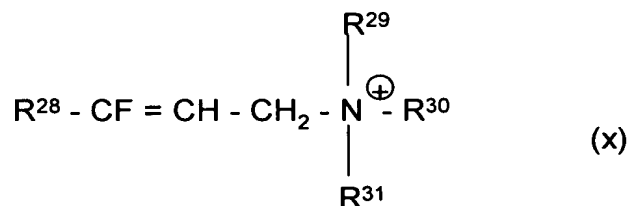
- 8) The method as claimed in claim 7, wherein the heterocyclic radical is pyridinium, pyridazinium, pyrimidinium, pyrazinium, purinium, tetraazaporphyrinium, piperidinium, morpholinium, tetrazonium, triaza-cyclononanium or tetraaza-cyclododecanium.

- 9) The method as claimed in claim 4, wherein the cationic metal complex is a metal carboxylate, metal salicylate, metal sulfonate, 1:1 metal-azo complex or a metal dithiocarbamate.

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- 10) The method as claimed in claim 9, wherein the metal is Al, Mg, Ca, Sr, Ba, TiO, VO, Cr, V, Ti, Zr, Sc, Mn, Fe, Co, Ni, Cu, Zn and ZrO, and the metal complex optionally contains one or more further ligands.
- 11) The method as claimed in claim 1, wherein the organic cation is a fluorinated ammonium ion of the formula (x)



- in which

R^{28} denotes perfluorinated alkyl having 5 to 11 carbon atoms and

R^{29} , R^{30} and R^{31} are identical or different and denote alkyl having 1 to 5 carbon atoms.

12) Salt-like structured silicate, in which the silicate is hectorite, beidellite, illite, muscovite, xantophyllite, margarite, sepiolite, saponite, mica, feldspar, nontronite, montmorillonite, smectite, bentonite, faujasite, zeolite A, X or Y, permutite, sasil or a combination thereof; and the cation is an ion of the formula (x) as claimed in claim 9.

13) A process for the preparation of a salt-like structured silicate as claimed in claim 12, which comprises combining the silicate and a salt of the cation of formula (x) in an aqueous medium.

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14) An electrophotographic toner, powder or powder coating comprising 30 to 99.99% by weight of a binder, 0.01 to 50% by weight, of at least one salt of ionic structured silicates in which the cation is NH_4^+ , H_3O^+ , an alkali metal, alkaline earth metal, earth metal or transition metal ion or a low molecular weight organic cation or a combination thereof and the anion is an island, cyclic, group, chain, ribbon, laminar or matrix silicate or a combination thereof, and optionally 0.001 to 50% by weight, of a coloring agent, in each case based on the total weight of the electrophotographic toner, powder or powder coating.

15) An electrophotographic toner, powder or powder coating as claimed in claim 14, comprising 40 to 99.5% by weight of a binder, 0.05 to 20% by weight of at least one salt of ionic structured silicates in which the cation is NH_4^+ , H_3O^+ , an alkali metal, alkaline earth metal, earth metal or transition metal ion or a low molecular weight organic cation or a combination thereof and the anion is an island, cyclic, group, chain, ribbon, laminar or matrix silicate or a combination thereof, and optionally 0.05 to 20% by weight of a coloring agent, in each case based on the total weight of the electrophotographic toner, powder or powder coating.

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